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OM nucleic - nucleic search, using sw model

Run on: September 23, 2003, 23:11:13 ; Search time 579 seconds

(without alignments)

9818.681 Million cell updates/sec

Title: US-09-856-327-1

Perfect score: 2106

Sequence: 1 atgcacatgtctctctca.....aaaaaaaaaaaaaaaaaaaaa 2106

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 252756 seqs, 1349719017 residues

Total number of hits satisfying chosen parameters: 5105512

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

N_Geneseq_19Jun03:*

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2: /SIDS1/gcgdata/geneseq/geneseq-nbml/NA1981.DAT:*

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24: /SIDS1/gcgdata/geneseq/geneseq-nbml/NA2002.DAT:*

25: /SIDS1/gcgdata/geneseq/geneseq-nbml/NA2003.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	2106	100.0	2106	AAF99980	Nucleotide sequenc
2	212.4	10.1	1869	AAAC87519	Trametes hirsuta p
3	212.4	10.1	1995	AAAC87518	Trametes hirsuta p
4	208.6	9.9	1869	AAAT34420	Pyranose oxidase e
5	175.6	8.3	1902	AA246411	Pleurotus cornucop
6	152.6	7.2	1701	AAV83626	Nucleic acid encod
7	129	6.1	1946	AAA71487	T. matsutake pyran
8	129	6.1	1946	AAA07403	Trichoderma derive

9	94.4	4.5	4639	22	AA545962	Human DNA encoding
10	94.4	4.5	4640	21	AA578483	Human PRO708 (UNQ3
11	94.4	4.5	4640	25	ACA57720	Human PRO708 cDNA.
12	94.4	4.5	4640	25	ABX98190	Human cDNA encodin
13	94.4	4.5	4640	25	ABX98692	Novel human secret
14	94.4	4.5	4640	25	ACA05737	Human secreted/tra
15	94.4	4.5	4640	25	ABX97781	Human PRO polynucl
16	94.4	4.5	4640	25	ABX78565	Human PRO polynucl
17	94.4	4.5	4640	25	ABX75578	Human cDNA encodin
18	94.4	4.5	4640	25	ABX76783	Human PRO polynucl
19	94.4	4.5	4640	25	ABX16623	Human cDNA encodin
20	94.4	4.5	4649	25	ABX92359	Human cDNA encoding
21	94.4	4.5	4650	20	AAZ33987	Human PRO708 nucle
22	93.8	4.5	552	23	ABV58513	Human prostate exp
23	92.6	4.4	269	23	ABV07492	Human prostate exp
24	92.2	4.4	488	23	ABV58909	Human prostate exp
25	91.8	4.4	2440	22	AAH34932	Human colon cancer
26	91.6	4.3	485	23	ABV55622	Human prostate exp
27	91.6	4.3	2323	19	AAV59524	Human prostate exp
28	91.6	4.3	2323	24	ABV73511	Human secreted pro
29	91.4	4.3	1480	22	AAAD07771	Human cDNA #1 for
30	91.2	4.3	2091	24	AAAL50827	Human secreted pro
31	91.2	4.3	2710	21	AAA54132	Human cancer statu
32	91.2	4.3	2964	24	ABK70015	Breast cancer prot
33	91.2	4.3	422	23	ABV54507	cDNA encoding huma
34	90.8	4.3	1327	24	AAAD24775	Human prostate exp
35	90.8	4.3	1565	25	ACC00654	Glycine max ankyri
36	90.6	4.3	1992	22	AAV72748	Human prostate can
37	90.6	4.3	2537	25	ABV76135	Argemone mexicana
38	90	4.3	198	23	ABV19260	Coxsackievirus rec
39	90	4.3	400	23	ABV58615	Human prostate exp
40	90	4.3	481	23	ABV56248	Human prostate exp
41	90	4.3	579	23	ABV57511	Human prostate exp
42	89.6	4.3	374	22	AAI91055	Human polynucleoti
43	89.6	4.3	1046	24	AA563134	Cell death protect
44	89.6	4.3	2260	22	AAI97914	Human neuroblastom
45	89.6	4.3	2260	22	AAI98068	Human neuroblastom

ALIGNMENTS

RESULT 1
AAF99980

ID AAF99980 standard; cDNA; 2106 BP.

XX

AC AAF99980;

XX

DT 20-JUL-2001 (first entry)

XX

DE Nucleotide sequence encoding Lyophyllum shimeji antibacterial protein.

XX Fungus; antibacterial; antibiotic; plant pathogen; bacterial infection;

XX Pyricularia oryzae; Rhizoctonia solani; rice pathogen; ss.

XX

OS Lyophyllum shimeji.

XX

FH Key Location/Qualifiers

CDS

FT 8..1864

FT /*tag= a

FT /product= "antibacterial protein"

XX

PN WO200121657-A1.

XX

PD 29-MAR-2001.

XX

PF 20-SEP-2000; 2000WO-JP06404.

XX

PR 21-SEP-1999; 99JP-0267238.

XX

PA (NISR) JAPAN TOBACCO INC.

PA (NORQ) SOC TECHNO-INNOVATION AGRIC FORESTY & FI.

Db	1741	GGGCTTGGCGAGAACCCGACACTTACGTGCGATGTGCCACGCTATCAGAGCGCGAGGAG	1800
Qy	1801	CATCATCAATACACTCAAGGGTGGGACTGACGGAAAAAATACAGCGGAGCATCGCAACCT	1860
Db	1801	CATCATCAATACACTCAAGGGTGGGACTGACGGAAAAAATACAGCGGAGCATCGCAACCT	1860
Qy	1861	TTGAGGAGGAGCAACAGCAGTGTAAACAACGCGTCAAGTGGCTACTTCAAGTTCGAATG	1920
Db	1861	TTGAGGAGGAGCAACAGCAGTGTAAACAACGCGTCAAGTGGCTACTTCAAGTTCGAATG	1920
Qy	1921	CATCTGGTCCCCCTACCATGTTGATGTGTACGATAGGCGTTGAAAGATTGTGTGTATTAC	1980
Db	1921	CATCTGGTCCCCCTACCATGTTGATGTGTACGATAGGCGTTGAAAGATTGTGTGTATTAC	1980
Qy	1981	TGAACCTGTACTTTGTCTGTAATAGTATGCGACTATGATCATGTTTAAAAAANAANAANA	2040
Db	1981	TGAACCTGTACTTTGTCTGTAATAGTATGCGACTATGATCATGTTTAAAAAANAANAANA	2040
Qy	2041	AAAAAANAANAANAANAANAANAANAANAANAANAANAANAANAANAANAANAANAANA	2100
Db	2041	AAAAAANAANAANAANAANAANAANAANAANAANAANAANAANAANAANAANAANAANA	2100
Qy	2101	AAAAAA 2106	
Db	2101	AAAAAA 2106	

RESULT 2	
XX AAC87519	
ID AAC87519 standard; DNA; 1869 BP.	
XX AC AAC87519;	
XX AC AAC87519;	
XX 13-MAR-2001 (first entry)	
XX	
XX Trametes hirsuta pyranose oxidase cDNA, SEQ ID NO:1 (version 2).	
DE	
XX Pyranose oxidase; expression construct; recombinant production;	
KW monosaccharide oxidation; 2-keto derivative;	
KW hydrogen peroxide production; ss.	
XX	
XX Trametes hirsuta.	
OS	
XX US6146865-A.	
PN	
XX 14-NOV-2000.	
PD	
XX 05-MAY-1999; 99US-0305381.	
XX	
XX 08-JUN-1998; 98DK-0000774.	
PR	
XX 10-JUN-1998; 98US-0088724.	
XX	
XX PA (NOVO) NOVO NORDISK AS.	
XX	
PI Schneider P, Christensen S, Lassen SF;	
XX	
PI WPI; 2001-049055/06.	
DR	
XX P-PSDB; AAB48832.	
XX	
XX Novel nucleic acid molecule encoding polypeptide having pyranose	
PPT oxidase activity used to design oligonucleotide probes to identify and	
PPT clone DNA encoding the polypeptide from different genera or species -	
XX	
XX Claim 2; Column 25-28; 20pp; English.	
XX	
XX The invention relates to nucleic acids (e.g., AAC87518, AAC87519) which	
CC encode Trametes hirsuta pyranose oxidase (AAB48832). The invention also	
CC relates to expression constructs, expression vectors and recombinant	
CC cells comprising pyranose oxidase nucleic acid sequences, and the	
CC recombinant production of Trametes hirsuta pyranose oxidase. Pyranose	
CC oxidase catalyses the oxidation of several monosaccharides in the	
CC pyranose form at position C2 to produce 2-keto derivatives with the	
CC release of hydrogen peroxide. Nucleic acids encoding Trametes hirsuta	

Db 1051 CCGTACCTGGGACCCACATCACCGAGCAGCGCTGCTTCTGCGACGCTCATGAGC 1110
QY 1160 CAGGAATTCGTGACAGCGTGGCGGAGTC----- 1190
Db 1111 ACGGAGCTCATCAACAGTGTCCCGGATATGACCAATGTGCGCAAGCCCGGCCACCGG 1170
QY 1191 -----CTTATGGACTGCCATGGTGGAAA 1213
Db 1171 GACTATAGCGTCACGTATACCCCGGGCCACCGACACAGACACCGGACTGGTGGAC 1230
QY 1214 GAAGCGGTGCTACATATATGCCAAGAACCGCAGATGACACTGCCATTCGTTCCGC 1273
Db 1231 GAGAAGGTGAAGAAGCACATGATGGACCAACAGGAGGACCGCTCCCGATCCGTTCCGAG 1290
QY 1274 GATCGGACCCCGGTACACACCCCATTTACAGAAACACCCCTGGCAGCAGATT 1333
Db 1291 GACCTGAGCCGCGGTACACAGCTGTTTTCAGGCAACGCCACCCATGGCACACCCAGATT 1350
QY 1334 CACCGCGATGCTTTTTCGTACGTCGCTCGGTCCTGAGTGGACTCTCGTGCATCGTC 1393
Db 1351 CACCGCGAGCCTTCAGCTACGGCCGCTGCGACGAGCATCGACTCGGGCTCATCGTC 1410
QY 1394 GACCTGCGGTGTTGGCGCAACCGACCGCTGAAGCAAAACACTTTTGTGTTTCCAGAAC 1453
Db 1411 GACTGGCGGTTTTCGGACGACCGACCGCCCAAGGAGGAGAACAGCTATGGTTCFCGGAC 1470
QY 1454 GATGTTCAAGCGGTACAGTATGCCGACCGCGCTTCAGATATCGACCCAGCACTCGG 1513
Db 1471 AGATCAGGAGCGGTACACCTCCGGCAGCGCGCTTCGACTTCCTGCTCCCGGGG-- 1528
QY 1514 TCAACAGTGAGAGGAAGAAATATGTCGGAAGTGGCGAGCAACTTGGGA 1573
Db 1529 ----GCCGGAAGCGGAGGACATGATGACCGACATGTGGTTCATGTCGGCGAAGATCGGT 1584
QY 1574 GGTATTTGCCAGCTCCCGCCGAGTTATGATGATGATGATGATGATGATGATGATGATG 1633
Db 1585 GATTCCTGCTGGGCTTCCACACAGTTCATGAGAGCGCGCTTGTCTGCTGACCTGCT 1644
QY 1634 GGGACTACTCGGATTTGCTGACAA-----GGCAACTACAGTGGCTGATACAACTCG 1687
Db 1645 GGGAGCAGCGATGGGTTTCGACGAGAGCGGACAAAGTGTGCTGCGACCGCACTCA 1704
QY 1688 CTGGTCTGGGATTTGCAATCTTTATGTTGAGGCAATGGCAACCATCAGGACGGGCTTC 1747
Db 1705 CGCGTCTGGCTTCAAGAACTCTTCTCGCGGCTGCGGGAACATCCCAACCGCTAC 1764
QY 1748 GCGAGAACCCGACACTTACGTGATGTCGACGCTATCAAGAGCGGAGGAGCATCATC 1807
Db 1765 GCCGGAACCCGAGCTCACCGCAATGTGCTTGGATCAAGAGCTGCGAGTACATCAAG 1824
QY 1808 AATACACTCAAG 1819
Db 1825 AAGAACTCGAG 1836

RESULT 3

AAC87518

ID AAC87518 standard; DNA; 1995 BP.

XX AAC87518;

AC AAC87518;

DT 13-MAR-2001 (first entry)

DE Trametes hirsuta pyranose oxidase cDNA, SEQ ID NO:1 (version 1).

KW Pyranose oxidase; expression construct; recombinant production;

KW monosaccharide oxidation; 2-keto derivative;

XX hydrogen peroxide production; ss.

OS Trametes hirsuta.

XX US6146865-A.

PN

XX 14-NOV-2000.
PD 05-MAY-1999; 99US-0305381.
XX 08-JUN-1998; 98DK-0000774.
PR 10-JUN-1998; 98US-0088724.
XX (NOVO) NOVO NORDISK AS.
PA
XX
PI Schneider P, Christensen S, Lassen SF;
XX WPI: 2001-049055/06.
DR P-PSDB; AAB48832.

XX Novel nucleic acid molecule encoding polypeptide having pyranose
PT oxidase activity used to design oligonucleotide probes to identify and
PT clone DNA encoding the polypeptide from different genera or species -
XX Claim 2; Fig 1; 20pp; English.

XX The invention relates to nucleic acids (e.g., AAC87518, AAC87519) which
CC encode Trametes hirsuta pyranose oxidase (AAB48832). The invention also
CC relates to expression constructs, expression vectors and recombinant
CC cells comprising pyranose oxidase nucleic acid sequences, and the
CC recombinant production of Trametes hirsuta pyranose oxidase. Pyranose
CC oxidase catalyses the oxidation of several monosaccharides in the
CC pyranose form at position C2 to produce 2-keto derivatives with the
CC release of hydrogen peroxide. Nucleic acids encoding Trametes hirsuta
CC pyranose oxidase may be used to produce the enzyme and to design
CC oligonucleotide probes to identify and clone genomic pyranose oxidase
CC cDNA or genomic DNA from different genera or species of microorganisms
CC (fungi or bacteria). The present sequence represents a cDNA encoding
CC pyranose oxidase from the fungus Trametes hirsuta.

CC Note: Both AAC87518 and AAC87519 are Trametes hirsuta pyranose oxidase
CC cDNA sequences which contain the entire open reading frame (ORF).
CC However, the two sequences have different stop codons - AAC87518 has an
CC opal stop codon while AAC87519 has an amber stop codon.

XX Sequence 1995 BP; 437 A; 649 C; 574 G; 335 T; 0 other;

Query Match 10.1%; Score 212.4; DB 22; Length 1995;
Best Local Similarity 51.1%; Pred. No. 3.2e-25;
Matches 841; Conservative 0; Mismatches 656; Indels 135; Gaps 10;

QY 269 GCCTACACACAGAGATGAATTCAGTTCAGAAAGATATTGACCGTTCTGCTCAATGTA 328
Db 288 GGTCTACACAAAGAGAACACCGTTCGAGTACCAGAAACATCGACAAATTCGTAATGTT 347

QY 329 ATCAAGGGAGCGTTACACAAAGTCTCTGTTCTGTCAGAACACGAGTGTGCTACACTT 388
Db 348 ATCAAGGGAGCGTTATGCGCGTCTGCTGCGGCTCAACACGATGCTGTGACACGCTA 407

QY 389 GATCCCGGAGCGTGGAGCGCGCCCTCGGAAAGTTTCAGGCATATCAAGCAATCTCT 448
Db 408 AGCCCGGCGTATGCGAGCTTCGACG-----TCTCTCGCGCAAGGGCGCAATCCA 461

QY 449 CACGAGCGGGAATTCGAGAACTGTCTCGGAGGCGCGTAAACGCTGGAGTCCGGCGCATG 508
Db 462 GAGCAAGACCGCTGCGCAACCTTAGTGGCCACCGCGGTCAACCGGCTGCTGCGCGCATG 521

QY 509 AGTACCCACTGGAGCTGCTCCACCGCCAGGATTCATCCACCCATGGAAGTCTCCCGGGC 568
Db 522 TCTACGCACTGGAGCTGCGGACCGCGCTTCGAGAAAGCTGCGAG----- 566

QY 569 ATCGGCGCTCGGAAGCTCAGTAACGACCGCGCAGGAGCAAGAGTGAACGAGCTT 628
Db 567 ---CGCCCGCTGCTCGTGAAGAACACTCCNAGCGGAGACGCGCGGAGTGGAGCAGGCTC 623

QY 629 TATTCCGAGCGCGAGCGTCTCATCGGAGCTTCCACCAAGGAATTCGACGAGTCAATCGG 688
Db 624 TACAAGAACCGCGAGTCTACTTCAAGACCGGCGACGACCCAGTTCGCGGAGTCCGATCCG 683

QY	689	CACACCTTGTTCGCGCTCTTTTGCAGAACCGGTACAAAGGATCGTCAACGATATCTTTTCG 748
DB	684	CACAACTCTGTGCTCAAGAACTCGAGGAGAGTACAAG---CGCTGCGGACTTCACG 740
QY	749	CCTCTCCGTTGGCATGCCACCGGTTGAAGAACCGCGGATACGTCGATGGCACTCA 808
DB	741	CAGATCCGCTCGGGGACGCGCCAGA-----GCCCGAGCTTCGTCGAGTGGAGCTCG 794
QY	809	GCAGAAATCTTTTCCACTCTATCTACAACGATGACAAGCAAGAAAGCTCTTTTACCCTG 868
DB	795	CGCACACCGGTGT-----CGATCTCGAGACCGCGCGCAACAGGACGCG 839
QY	869	CTGACGAACCATCGCTGCACACGACTGGCGCTTACGGGCGGGTATGAGAAGAAATGGC 928
DB	840	CCGAAGCAGCGCTTCAACCTCTTCCCGCGCGTCCGCTGCACGAACGTGAGGCGCGATAAC 899
QY	929	GCTCCGAGGTCAAGAACTACTTCTGGCCACACAGGAATCCTAGTTCCGAGCTGGACAGCTAT 988
DB	900	CGCAACTCGGATCTGATAGGCTCGATGTCGCGACCTCCACGGGGCAAGAGCATCACC 959
QY	989	ATCATGCGGAAGTATATGTACTGGCGTCGGGAGCGATCGGCAACCCACACAGATTCCTAT 1048
DB	960	ATCAAGCCCAAGGTGATACCTCTCACCGCGCGCGTCCACACGCGCAGCTCTCGCG 1019
QY	1049	AATCTGGGCTT-----CTCTGGGCTACAGGTACGCCACGCAATGAC---TGCTTGATC 1099
DB	1020	GCCTCTGGATTGGGCGACGTGGGTCTGCGGACCGCCGCCAAGCGCTGCGGTCTCTGCTG 1079
QY	1100	CCCAACTCTGGGAGGTACATCACGGACACCGATGCACTTTTGCCAGATAGTCTTGAGG 1159
DB	1080	CCGTACTCTGGGACCCACATCACCGACGACGCTGCTCTCTGCGACGCGTCATGAGC 1139
QY	1160	CAGGAATTCTGCACACGCTGCGCGAGGATC----- 1190
DB	1140	ACGGAGCTCATCAACAGTGTCAACCGCGATATGACCATTTCGCGCAAGCCCGCCACCG 1199
QY	1191	-----CTTATGGACTGCCATGGTGAAA 1213
DB	1200	GACTATAGCTACGTATATACCCGGGAACCCGAAACAAGCACCGGACTGGTGAAC 1259
QY	1214	GAAGCCGTTGCTCAACATATTGCCAAGAACCCGACAGATGCACTGCCATTCGTTCCCG 1273
DB	1260	GAGAGGTGAAGAAGCATGATGGACCACAGAGGACCGCTCCCGATTCGCCGTTGCGAG 1319
QY	1274	GATCCGGAACCCAGGTAAACACCCCATTTACAGAAGAACACCCCTGGCACACGCAGATT 1333
DB	1320	GACCTTGAGCGCAGGTCAACACGCTGTTTTCAGGCAACGACCCATCGGCACACCCAGATT 1379
QY	1334	CACCGCGATGCTTTTTCGTACGTTGCGTCCGCTCTGAGGTGGAATCTCGTGTCACTGTC 1393
DB	1380	CACCGCGACGCTTACGTTACGCGCGCGTGCACGAGAGCATGCACTCGGGCTCATCTGTC 1439
QY	1394	GACCTGCGCTGTTTGGCGCAACCGACCTGAAGCAACAACTTTTGGTTTTCAGAAC 1453
DB	1440	GATGCGGTTCTTCGAGCGCACCGACCCCAAGAGGAGAAAGATATGGTTCTCGGAC 1499
QY	1454	GATGTTCAAGACGGGTACAGTATGCGCGAGCCGACGTTTACAGATATCGACCCAGCTGCG 1513
DB	1500	AAGATCAGGACCGGTACAACTTCGCGACCGCACGCTTCGACTTCGCTTCCCGGGG--- 1557
QY	1514	TCAAACGTGAGCAAGGAAATGATGGCCGATATGTGCGAAGTGGCGAGCAACTTGGGA 1573
DB	1558	-----GCCCGGAAGCGGAGCATGATGACCGACATGTGGTCTATGTCGCGGAAGATCGGT 1613
QY	1574	GGTATTTCGCCAGCTCCCGCCGAGTTTATGGATCCAGGCGCTTCGACTTCATCTTTGGG 1633
DB	1614	GGATTCTGCTGGGTCTTACCACAGTTTCATGGAGCCCGGCTTGTCTCTGCACCTTGT 1673
QY	1634	GGGACTACTCGCATTTGGCTTCGACAA-----GGCAACTACAGTGGCTGATACAACTCG 1687
DB	1674	GGGACCGCGATGGGCTTCGACGAGAGAGGCGGACAGTGTGGCTCGACACCGACTCA 1733
QY	1688	CTGGCTGGGACTTTGCCAATCTTTATGTTGCAAGCAATGGACCAATCAGGACGGGCTTC 1747

Db	1734	CGCGTCTTCGGCTTC	AAGAACCTCT	CCTCGCGGCT	TGCGGGAACAT	CCCAACCGCGTAC	1793
QY	1748	GGCGAGAACCCGACAC	TACGTCTGAT	TGCGCACGCTAT	CAAGAGCGCGAGGAGCAT	CATC	1807
Db	1794	GC CGGAACCGAGCGT	CACCGAATCT	CGTTCGGATCA	AAGAGCTCGGAGTACATCAAG	1853	
QY	1808	AATACACTCAAG	1819				
Db	1854	AAGAACTTCGAG	1865				

RESULT 4

AAT34420
 ID AAT34420 standard; cDNA; 1869 BP.

XX
AC AAT34420:

DT 27-NOV-1996 (first entry)

DE pyranose oxidase encoding sequence.

Pyranose oxidase; glucose; oxidation; glucanose; assay; diabetes;
KW
marker; diagnosis; 1,5-anhydro-D-sorbitol; ss.
KW

OS *Coriolus versicolor*.

PN DE19545780-A1.

13-JUN-1996

PF 07-DEC-1995; 95DE-1045780.

PR 24-MAY-1995; 95JP-0124835.

XX
XX

SECRET

NOFORN

XX
XX
CONFIDENTIAL (U)

PI Suzuki M;
PI Suzuki M;
PI Suzuki M;

DR WPI; 1996-278990/29.

[illegible]

1,5-anhydro-D-sorbitol used as marker for diabetes diagnosis

PS Claim 1; page 10-13; 22pp; German.

CC The present sequence encodes a protein isolated from *Corioliolus versicolor*,
CC which has the enzyme activity of pyranose oxidase (PO). The PO oxidises
CC

CC glucose to glucosone and has an optimum pH of 7-7.5. It has a mol. wt. 230000 (determined by gel filtration) and is stable at around 50deg.C.

CC PO can be used for measurement of glucose in, e.g. foods or body fluids.

CC or 1,5-anhydro-D-sorbitol which is an important marker used in the diagnosis of diabetes.

Sequence 1869 bp: 408 A: 601 C: 521 G: 339 T: 0 other: xx

Query Match 9.98: Score 208.6: DB 17: Length 1869:

Matches 856; Conservative 0; Mismatches 714; Indels 129; Gaps 9
 best Local Similarity 50.4%; Pred. NO. 1.3e-24;

190 GAGCTCCGCGAAGCCCGGTATACAGGCTCCGCAATGCTTCGACATCTGCTGGC
 200 GAGATCGGAGCTGCTGATAGCTTCTACCGTGTTAATGCCGAAGAAGAACTGCAGTTCCC
 250

Qy 250 TAGGTTCTCTGGCTACCAACAAGAAGATGAATCGAGTTCACAGAAAGATATTGACCGGCTC 31

DB 250 CTGAAGATCGGTGCCCAACAAGAAGAACACCGTCGAATACCAAGAGAACATTGCACAAG

Db 310 GTGAAGCTATTAGAGGCAATTGATGCTGTGTTCCGTTCCGTCATACCCCTCGTGATC 369
QY 380 CCTACACTTATCCCGAGCGTGGAGCGCCGCCCTGGAAGTTTACGCCATATGAACGGT 439
Db 370 GACAGCTAGCCCGAGCGTGTGCAAGCTTCATCG-----TTCCTCGTCCCAATGGC 423
QY 440 AAAAATCCTCACCAGCGGGAATTCGAGAACTTGTCTGCGGAGGCGGTAAAGCGTGGAGTC 499
Db 424 TCGNACCCAGAGGACCGCGTTTCGTAACCTCAGTGGTCAGCGGCTCAGCGTGTGCTC 483
QY 500 GGGCGCATAGTACCCACTGGAGTGTCTCCACGCCACGGATTATCCACCCATGGAAAGT 559
Db 484 GGAGGATGTCACGCACTGGACATGCGGACACCGCGCTTGA-----527
QY 560 CTCGCCGGCATCGCGCTCGGAAGCTCAGTAACGACCGCGCAGAGGACGACAAAGAGTGG 619
Db 528 --CCGGGAGCGCGCCGTTGCTCGTGAAGGAGCAGCAGGACGCTGACGACGCGAGTGG 585
QY 620 AACGAGCTTATTCCGAGCGGAGCGTCTCATCGGACTTCCACCAAGGNAATTCGACGAG 679
Db 586 GACCGGCTGTACACCAAGCGCGAGTCTACTTCAAGACCGGAGCAGCAGCTGACGACGCGAGTGG 645
QY 680 TCAATTGCGCACACCTTGTCTGCGCTCTTTGCAAGACGCGGTACAAAGGATCGTCAACGT 739
Db 646 TCGATCGGCACAACTCTGCTCAACAGCTCGCGGAGGATACAAAGTTCAGCGCGAC 705
QY 740 ATCTTTGCGCCCTCTCCGTTGGATGCCACCGGTTGAAGACGCGCGGAAATACGTCGAA 799
Db 706 TTCCAGCAGATCCCCCTCGCGCAACGCGTCG-----CAGTCCGACCTTCGTCGAG 756
QY 800 TGGCACTCAGCAGAAATCTTTTCCACTTATCTACACGATGACAGCAAGAGAGCTC 859
Db 757 TGGAGCTCGGCAACACCGTGT-----CGACCTCCAGAACAGGCGCGAAC 801
QY 860 TTATCCCTGCTCAGCAACCATCGCTGCACAGCTGGCGCTTACGGCGGGTATGAGAAG 919
Db 802 ACGGACGCGCCGAATGAGCGCTTCAACCTTTCGCCGCGTTCATGTGAGCGCGTGTG 861
QY 920 AAGATTGGCGTTCGCGAGTCAAGATCTACTGGCCACGAGGAATCTTAGTTCCGAGTGT 979
Db 862 CCACACAGCTGCACTCGAGATCGAGATGTGCACATCCACGACCTCATCTCCGCGAC 921
QY 980 GACAGCTATATCATGCGAAGGTATATGTACTGGCGTGGGAGCGATCGGCAACCCACAG 1039
Db 922 CGCTTCGAAATCAGACAGACGTGTGCTTCTTACGCGGGCGGTCCACACCGCGAG 981
QY 1040 ATTCTCTATACTCGGGT-----CTCTGGGCTACAGGTCAAGCCACGCAATGACTCG 1093
Db 982 CTTCCTGTAACCTCTGGCTTTGGACAGCTGGGCGCGCGACCCCGCAACCCCGCGAG 1041
QY 1094 TTGATCCCAACCTGGGAGGTACATCAGGAGGAGCGGATGGCATTTTGGCAGATGTC 1153
Db 1042 TTGCTGCGCTCCCTCGAAGTATACATCAGGAGAGTGTCTGCTTCTGCGCAGACCGTG 1101
QY 1154 TTGAGGAGGAATTCGTCGACAGCTG-----CGGAGCATCCTTATGCACTGCC-----1203
Db 1102 ATGACCCGAGCTCATCGACGCTGAAGTCCGACATGATCATCAGGGGCAACCTGGC 1161
QY 1204 -----ATGG 1207
Db 1162 GATCTGGGTTACAGGTCACGTACAGCGCGCGGAGACCAACAGCAGCCCGGACTGG 1221
QY 1208 TGGAAAGAGCGGTGCTCAACATATTGCCAAGAACCGGACAGATGCACTGCCATTCGG 1267,
Db 1222 TGGAAAGAAAGGTGAAGAACCCACATATGACGACCCAGGAGGACCGCTTCCCAATCCCG 1281
QY 1268 TTCCGCGATCGGAAACCCAGGTAAACACCCATTACAGAGAACACACCCCTGGCACAG 1327
Db 1282 TTCAGAGACCCGAGCGAGGTCAACACCTTGTTCAGGCATCGCAGCCGTCGACACT 1341
QY 1328 CAGATTACCGCGATGCTTTTTCGTAGCGTGCCTCGTCTCTAGGTGGACTCTCGGTC 1387
|||||

Db 1342 CAGATTACCGCGGATCGTTAGTTACGGCGCGTGTGAGCAAAACCATCGACTCAGCTC 1401
QY 1388 ATGCTGACCTGGCGTGTGTTGGCGCAACCGACCTGAAGCAAAACACTTTTGGTTTTC 1447
Db 1402 ATGCTGACCTGGCGTGTGTTGGCGCGGAGGAGCAAGGAGGAGAACAGCTCTGGTTC 1461
QY 1448 CAGAACATGTTCAAGAGGATACAGTATGCCGAGCGAGCTTTCAGATATCGACCCAGC 1507
Db 1462 TCGGACAAAATACGGACAGTACACATGCGCGAGCGAGCTTCGACTTCGCTCCG 1521
QY 1508 ACTG---CGTCAACGTCGAGGCAAGAAATGATGCCCATATGTGCGAGTGGCGAGC 1564
Db 1522 GCGGCGCGACGAGCAAGGAGGCGGAGGACATGATACCGCATATGTGCGTATGTCGCG 1581
QY 1565 AACTTGGGAGTATTATTGCCACGCTCCCGCGAGTTTATGGATCCAGGCTTGCATTT 1624
Db 1582 AAGATTGGTGGCTTCTGCCCGCTCCCTCCGCAATTCATGAGCCCGGCTTGTGCTT 1641
QY 1625 CATCTTGGGGGACTACTTCGATCTGGCTTCGAC-----AAGCAACTACAGTGGCTGAT 1678
Db 1642 CACCTGGTGTACGACCGCATGGCTTCGAGGAGGAGGACAAAGTGTGGTCAAC 1701
QY 1679 AACAACTCGCTGCTGGGACTTTGCCAATCTTTATGTTGTCAGCAATGSCACCATCAGG 1738
Db 1702 ACGGACTCGCGCTGTGGCTTCAAGAACCTTCTCTCGTGGTGGCGGAAACATTTCC 1761
QY 1739 ACGGGCTTGGCGGAGAACCGACACTTACGTCATGTGCCACGCTATCAAGAGCGCGAGG 1798
Db 1762 ACGCGTACGCGGAAACCGAGCTCACCGCAATGTCCGTAATGTCCGTCGATCAAGATGTCGGAG 1821
QY 1799 AGATCATCAATACATCA 1817
Db 1822 TATCAAGAACAATTCA 1840

RESULT 5

AZ46411
ID AZ46411 standard; DNA; 1902 BP.

XX AZ46411;

XX 07-MAR-2000 (first entry)

XX Pleurotus cornuoplae antitumour protein coding sequence.

XX Antitumour; cancer; tumour; treatment; expression; tumour suppressor;

KW p53; pBR; ss.

XX Pleurotus cornuoplae.

XX JP11315096-A.

XX 16-NOV-1999.

PF 07-AUG-1998; 98JP-0236349.

XX 08-AUG-1997; 97JP-0215311.

PR 02-MAR-1998; 98JP-0066176.

XX (NEWF-) NEW FOOD CREATION GIJUTSU KENKYU KUMIAI.

XX WPI; 2000-058170/05.

DR P-ESDB; AAY52700.

XX An antitumour protein derived from Pleurotus cornuoplae and its gene -
useful for treatment of cancer including those caused by abnormal
expression of cancer inhibitory gene (e.g. p53 and pBR)

XX Claim 9; Page 15-16; 23pp; Japanese.

CC The invention relates to a novel antitumour protein extracted from
fruiting bodies of the fungus Pleurotus cornuoplae. The protein and
nucleotides encoding it are useful for the treatment of cancer,

PR 25-APR-2000; 2000US-199654P.
 PR 03-MAY-2000; 2000US-201516P.
 PR 17-MAY-2000; 2000WO-US13705.
 PR 22-MAY-2000; 2000WO-US14042.
 PR 30-MAY-2000; 2000WO-US14941.
 PR 02-JUN-2000; 2000WO-US15264.
 PR 05-JUN-2000; 2000US-209832P.
 PR 28-JUL-2000; 2000WO-US20710.
 PR 22-AUG-2000; 2000US-0644848.
 PR 24-AUG-2000; 2000WO-US23328.
 PR 08-NOV-2000; 2000WO-US30952.
 PR 01-DEC-2000; 2000WO-US32678.
 PR 20-DEC-2000; 2000WO-US34956.
 XX (GETH) GENENTECH INC.
 PA Baker KP, Chen J, Desnoyers L, Goddard A, Godowski PJ, Gurney AL;
 PI Pan J, Smith V, Watanabe CK, Wood WI, Zhang Z;
 XX
 DR WPI: 2001-602746/68.
 P-PSDB; AAU29061.
 XX
 XX Novel nucleic acids encoding PRO polypeptides, used to diagnose the
 PT presence of tumours, such as prostate and breast tumours, in mammals and
 PT to screen for modulators of the compounds
 XX
 PS Claim 2; Fig 75; 774pp; English.
 XX
 CC Sequences AAS45925-AAS46231 represent DNA molecules encoding and PCR
 CC primers for PRO polypeptides of the invention. The sequences of the
 CC invention can be used to detect the presence of a tumour in a mammal by
 CC comparing the level of expression of a PRO polypeptide in a test sample
 CC of cells from the animal and a control sample of normal cells, whereby a
 CC higher level of expression in the test sample indicates the presence of a
 CC tumour in the mammal. Mammals include dogs, cats, cattle, horses, sheep,
 CC pigs, goats and rabbits but are preferably human. The polypeptides can be
 CC used to stimulate tumour necrosis factor (TNF) alpha release from human
 CC blood, when contacted with it. A specific polypeptide can be used to
 CC stimulate the proliferation or differentiation of chondrocyte cells. The
 CC PRO proteins can be used to determine the presence of tumours and also
 CC susceptibility to tumour development, particularly adrenal, lung, colon,
 CC breast, prostate, rectal, cervical, or liver tumours, in mammalian
 CC subjects. The oligonucleotide probes specific for the PRO nucleic acids
 CC can be used for genetic analysis of individuals with genetic disorders.
 XX
 SQ Sequence 4639 BP; 1426 A; 956 C; 1025 G; 1232 T; 0 other;
 Query Match 4.5%; Score 94.4; DB 22; Length 4639;
 Best Local Similarity 72.9%; Pred. No. 1.9e-06;
 Matches 137; Conservative 0; Mismatches 46; Indels 5; Gaps 1;
 QY 1924 TCTGGTCCCTACCATGTTGATGTGATGATGCGGTGAAAGATTTTGTGTTACTGA 1983
 Db 4423 TATGGCTCATTTTATTTATAGTGTAAAGTTGTATTTCTTAAAGTTTGTGTTCTCGA 4482
 QY 1984 ACCTGTACTTTGCTGA-----ATAGTTATGGCAGCATGATTCATGTTTAAAAA 2038
 Db 4483 CAGTATCTTTTAAATGAGTCTTAAAAATAAGCATATTTGTCATGTTTAAAAA 4542
 QY 2039 AAAAAA 2106
 Db 4543 AAAAAA 4610
 QY 2099 AAAAAA 2106
 Db 4603 AAAAAA 4610

RESULT 10
 AAC78483
 ID AAC78483 standard; cDNA; 4640 BP.
 XX
 AC AAC78483;

XX 08-FEB-2001 (first entry)
 DT Human PRO708 (UNQ372) nucleotide sequence SEQ ID NO:113.
 DE Human; secreted protein; transmembrane protein; PRO; EST; cytosolic;
 XX expressed sequence tag; detection; cancer; ss.
 KW Homo sapiens.
 OS WO2000053756-A2.
 PN 14-SEP-2000.
 XX 18-FEB-2000; 2000WO-US04341.
 PF 08-MAR-1999; 99WO-US05028.
 XX 12-MAR-1999; 99US-0123957.
 PR 29-MAR-1999; 99US-0126773.
 PR 21-APR-1999; 99US-0130232.
 PR 28-APR-1999; 99US-0131445.
 PR 14-MAY-1999; 99US-0134287.
 PR 23-JUN-1999; 99US-0141037.
 PR 26-JUL-1999; 99US-0145698.
 PR 29-OCT-1999; 99US-0162506.
 PR 30-NOV-1999; 99WO-US28313.
 PR 02-DEC-1999; 99WO-US28551.
 PR 16-DEC-1999; 99WO-US30095.
 PR 30-DEC-1999; 99WO-US31243.
 PR 05-JAN-2000; 99WO-US31274.
 PR 06-JAN-2000; 2000WO-US00219.
 PR 06-JAN-2000; 2000WO-US00277.
 XX (GETH) GENENTECH INC.
 PA Ashkenazi AJ, Baker KP, Botstein D, Desnoyers L, Eaton DL;
 PI Ferrara N, Filvaroff E, Fong S, Gao W, Gerber H, Gerritsen ME;
 PI Goddard A, Godowski PJ, Grimaldi CJ, Gurney AL, Hillan KJ;
 PI Kijavini IJ, Kuo SS, Napier MA, Pan J, Paoni NF, Roy MA;
 PI Shelton DL, Stewart TA, Tumas D, Williams PM, Wood WI;
 XX WPI: 2000-611443/58.
 DR P-PSDB; AAB44257.
 XX Novel PRO polypeptides and polynucleotides used in detection methods,
 PT to target bioactive molecules to specific cells, and to modulate
 PT cellular activities
 XX
 PS Claim 2; Fig 42; 636pp; English.
 XX AAC78458 to AAC78599 represent polynucleotide and EST (expressed
 CC sequence tag) sequences which encode secreted or transmembrane PRO
 CC polypeptides. The PRO polynucleotides and polypeptides have cytostatic
 CC activity. The polynucleotides and polypeptides can be used for detecting
 CC the presence of PRO polypeptides in samples, for linking bioactive
 CC molecules to cells and for modulating biological activities of cells,
 CC using the polypeptides for specific targeting. The polypeptide targeting
 CC can be used to kill the target cells, e.g. for the treatment of cancers.
 CC The polypeptide pairs provide specific targeting of bioactive molecules
 CC to cells. AAC78600 to AAC78987 represent PCR primers and probes used in
 CC the isolation of the PRO polynucleotide sequences.
 XX
 SQ Sequence 4640 BP; 1427 A; 955 C; 1026 G; 1232 T; 0 other;
 Query Match 4.5%; Score 94.4; DB 21; Length 4640;
 Best Local Similarity 72.9%; Pred. No. 1.9e-06;
 Matches 137; Conservative 0; Mismatches 46; Indels 5; Gaps 1;
 QY 1924 TCTGGTCCCTACCATGTTGATGTGATGATGCGGTGAAAGATTTTGTGTTACTGA 1983
 Db 4423 TATGGCTCATTTTATTTATAGTGTAAAGTTGTATTTCTTAAAGTTTGTGTTCTCGA 4482

Qy	1984	ACCTGTACTTTGTCTGA-----ATAGTTATGGCAGCTATGATTCATGTTTAAAAA	2038
Db	4483	CAGTATCTTTAAATGAGTCTTAAAAATAAAGGCATATTGTTTCATGTTTAAAAA	4542
Qy	2039	AAA	2098
Db	4543	AAA	4602
Qy	2099	AAAAAAAA 2106	
Db	4603	AAAAAAAA 4610	
RESULT 11			
ACAS7720			
ID	ACA57720 standard; cDNA; 4640 BP.		
XX	ACA57720;		
XX	XX		
DT	10-JUN-2003 (first entry)		
XX	XX		
DE	Human PRO708 cDNA.		
XX	XX		
KW	* Human; PRO; secreted; transmembrane; cytostatic; TNF-alpha; blood; gene;		
KW	tumour necrosis factor alpha release; chondrocyte cell; proliferation;		
KW	differentiation; tumour; gene therapy; ss.		
XX	XX		
OS	Homo sapiens.		
XX	XX		
PN	US2003036143-A1.		
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PD	20-FEB-2003.		
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PF	02-JUL-2002; 2002US-0187600.		
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PR	16-SEP-1998; 98WO-US19330.		
PR	07-OCT-1998; 98WO-US21141.		
PR	01-DEC-1998; 98WO-US25108.		
PR	08-MAR-1999; 99WO-US05028.		
PR	14-MAY-1999; 99WO-US10733.		
PR	02-JUN-1999; 99WO-US12252.		
PR	01-SEP-1999; 99WO-US20111.		
PR	15-SEP-1999; 99WO-US21090.		
PR	01-DEC-1999; 99WO-US28301.		
PR	02-DEC-1999; 99WO-US28551.		
PR	30-DEC-1999; 99WO-US31274.		
PR	05-JAN-2000; 2000WO-US00219.		
PR	18-FEB-2000; 2000WO-US04341.		
PR	18-FEB-2000; 2000WO-US04342.		
PR	22-FEB-2000; 2000WO-US04414.		
PR	24-FEB-2000; 2000WO-US05004.		
PR	01-MAR-2000; 2000WO-US05601.		
PR	02-MAR-2000; 2000WO-US05841.		
PR	15-MAR-2000; 2000WO-US06884.		
PR	30-MAR-2000; 2000WO-US08439.		
PR	17-MAY-2000; 2000WO-US13705.		
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PR	02-JUN-2000; 2000WO-US15264.		
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PR	24-AUG-2000; 2000WO-US23328.		
PR	08-NOV-2000; 2000WO-US30952.		
PR	01-DEC-2000; 2000WO-US32678.		
PR	20-DEC-2000; 2000WO-US34956.		
PR	28-FEB-2001; 2001WO-US06520.		
PR	01-JUN-2001; 2001WO-US17800.		
PR	20-JUN-2001; 2001WO-US19692.		
PR	29-JUN-2001; 2001WO-US21066.		
PR	09-JUL-2001; 2001WO-US21735.		
PR	29-AUG-2001; 2001WO-US27099.		
PR	18-SEP-1997; 97US-059263P.		
PR	18-SEP-1997; 97US-059266P.		

PR 28-FEB-2001; 2001WO-US06520. 98US-088167P.
PR 01-JUN-2001; 2001WO-US17800. 98US-088202P.
PR 20-JUN-2001; 2001WO-US19692. 98US-088212P.
PR 29-JUN-2001; 2001WO-US21066. 98US-088217P.
PR 09-JUL-2001; 2001WO-US21735. 98US-088655P.
PR 29-AUG-2001; 2001WO-US27099. 98US-088722P.
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PR 28-MAY-1998; 98US-087098P. 98US-097022P.
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PR 04-AUG-1998; 98US-095282P. 98US-098893P.
PR 10-AUG-1998; 98US-095998P. 98US-098894P.
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PR 01-SEP-1998; 98US-098723P. 98US-098909P.
PR 02-SEP-1998; 98US-098803P. 98US-098910P.
PR 02-SEP-1998; 98US-098821P. 98US-098911P.
PR 02-SEP-1998; 98US-098843P. 98US-098912P.

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PR 09-SEP-1998; 98US-099602P.
PR 10-SEP-1998; 98US-099741P.

Query Match          4.5%; Score 94.4; DB 25; Length 4640;
Best Local Similarity 72.9%; Pred. No. 1.9e-06;
Matches 137; Conservative 0; Mismatches 46; Indels 5; Gaps 1;

QY 1924 TCTGGTCCCTACCATGTTGATGTGACGATAGCGGTGAAGATTTGTGTATTACTGA 1983
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Db 4423 TATGGCTCCATTTATTTATAGTGAAGTTGTATTTCTTAAGTTTGTGTTTCTCGA 4482
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QY 1984 ACCTGTACTTGTCTGA-----ATAGTTATGGCAGTATGATTTTAAAAA 2038
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RESULT 13
ABX98692
ID ABX98692 standard; cdNA; 4640 BP.
XX
AC ABX98692;
XX
DT 20-MAY-2003 (first entry)
XX
DE Novel human secreted and transmembrane protein PRO708 cDNA.
XX
KW Human; secreted protein; transmembrane protein; cytosolic;
KW gene therapy; TNF-Agonist-Alpha; chondrocyte stimulator; tumour;
KW adrenal tumour; lung tumour; colon tumour; breast tumour;
KW prostate tumour; rectal tumour; cervical tumour; liver tumour;
KW gene; ss.
XX
OS Homo sapiens.
XX
PN US2003036157-A1.
XX
PD 20-FEB-2003.
XX
PF 02-JUL-2002; 2002US-0188769.
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PR 16-SEP-1998; 98WO-US19330.
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PR 01-DEC-1998; 98WO-US25108.
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PR 15-SEP-1999; 99WO-US21090.
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PR 30-DEC-1999; 99WO-US31274.
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PR 30-MAY-2000; 2000WO-US14941.
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PR 28-JUL-2000; 2000WO-US20710.
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PR 20-DEC-2000; 2000WO-US34956.
PR 28-FEB-2001; 2001WO-US06520.
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PR 29-AUG-2001; 2001WO-US27099.
PR 26-JUN-1998; 98US-0105413.
PR 07-OCT-1998; 98US-0168978.
PR 06-NOV-1998; 98US-0187368.
PR 07-DEC-1998; 98US-0204054.
PR 03-MAR-1999; 99US-0254311.
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PR 30-JUL-2001; 2001US-0918585.
PR 06-AUG-2001; 2001US-0924419.
PR 13-AUG-2001; 2001US-0929404.
PR 16-AUG-2001; 2001US-0931836.
PR 28-AUG-2001; 2001US-0941992.
PR 04-SEP-2001; 2001US-0946374.
PR 15-JAN-2002; 2002US-0052586.
XX

(GETH) GENENTECH INC.

Baker KP, Chen J, Desnoyers L, Goddard A, Godowski PJ, Gurney AL;
Pan J, Smith V, Watanabe CK, Wood WI, Zhang Z;

WPI; 2003-332039/31.
P-PSDB; ABU67443.

PT New secreted and transmembrane PRO polypeptides and nucleic acids,
PT useful in gene therapy, in chromosome and gene mapping, as chromosome
PT markers, in tissue typing, and in chromosome identification.

XX Claim 2; Fig 75; 706pp; English.

XX The invention discloses human nucleic acids encoding secreted and
XX transmembrane (PRO) polypeptides. Also disclosed is an antibody that
XX specifically binds to the PRO polypeptide, a method for stimulating the
XX release of tumour necrosis factor alpha (TNF-alpha) from human blood by
XX contacting the blood a PRO polypeptide, a method for stimulating the
XX proliferation or differentiation of chondrocyte cells by contacting the
XX cells with a PRO polypeptide, a method for detecting the presence of a
XX tumour in a mammal and an oligonucleotide probe derived from any of the
XX PRO nucleotide sequences. The nucleotide sequences are useful as probes,
XX in chromosome and gene mapping, in generating antisense RNA and DNA, in
XX preparing PRO polypeptides by recombinant techniques and in gene therapy
XX (e.g. for replacement of defective gene). The PRO polypeptides are useful
XX as molecular weight markers for protein electrophoresis purposes, for
XX chromosome identification, as chromosome markers, as therapeutic agents,
XX for stimulating the release of TNF-alpha from human blood, for

CC stimulating the proliferation or differentiation of chondrocytes and
CC detecting the presence of a tumour. The PRO polypeptides and nucleic
CC acids may also be used diagnostically for tissue typing. The sequences
CC presented in ACA05700-ACA06004 are the cDNAs encoding the PRO
CC polypeptides of the invention.

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SQ Sequence 4640 BP; 1427 A; 955 C; 1026 G; 1232 T; 0 other;

Query Match 4.5%; Score 94.4; DB 25; Length 4640;
Best Local Similarity 72.9%; Pred. No. 1.9e-06;
Matches 137; Conservative 0; Mismatches 46; Indels 5; Gaps 1;

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Db 4483 CAGTATCTTTTAAATGAGTCTTAAAAATAAGGCATATTTGTTCTCTTAAAAA 4542
QY 2039 AAAAAA 2098
Db 4543 AAAAAA 2098
QY 2099 AAAAAA 2106
Db 4603 AAAAAA 4610

RESULT 15

ABX97781
ID ABX97781 standard; cDNA; 4640 BP.

XX
AC ABX97781;

DT 16-MAY-2003 (first entry)

DE Human PRO polynucleotide #38.

XX Human; PRO; gene; ss; cytostatic; chromosome mapping; gene mapping;
KW protein electrophoresis; tumour necrosis factor-alpha; TNF-alpha; blood;
KW chondrocyte differentiation; chondrocyte proliferation; tumour.

XX Homo sapiens.

PN US2003032102-A1.

PD 13-FEB-2003.

PF 17-JUN-2002; 2002US-0173697.

XX 16-SEP-1998; 98WO-US19330.

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PR 01-DEC-1998; 98WO-US25108.

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PR 02-JUN-1999; 99WO-US12252.

PR 01-SEP-1999; 99WO-US20111.

PR 15-SEP-1999; 99WO-US21090.

PR 01-DEC-1999; 99WO-US28301.

PR 02-DEC-1999; 99WO-US28551.

PR 30-DEC-1999; 99WO-US31274.

PR 05-JAN-2000; 2000WO-US00219.

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PR 24-FEB-2000; 2000WO-US05004.

PR 01-MAR-2000; 2000WO-US05601.

PR 02-MAR-2000; 2000WO-US05841.

PR 15-MAR-2000; 2000WO-US06884.

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Search completed: September 24, 2003, 02:08:55
Job time : 593 secs